



Receipt

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PATENT

Case Docket No. TETRAGN.002A

Date: December 23, 1999

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

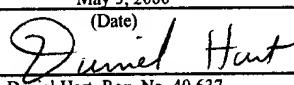
Applicant(s) : Irena N. Merenkova

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for Patents, Washington, D.C. 20231, on

Appl. No. : 09/471,703

May 5, 2000

(Date)

  
Daniel Hart

Daniel Hart, Reg. No. 40,637

Filed : December 23, 1999

For : ANALYSIS OF  
NUCLEOTIDE  
POLYMORPHISMS AT A  
SITE

Examiner : Unknown

Group Art Unit : 1643

**TRANSMITTAL LETTER**

**ASSISTANT COMMISSIONER FOR PATENTS**

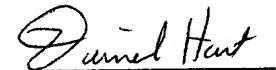
**WASHINGTON, D.C. 20231**

ATTENTION: Office of Initial Patent Examination's Customer Service Center

Dear Sir:

Enclosed for filing in the above-identified application are:

- (X) Request For Correction Of Filing Receipt.
- (X) Copy of the Official Filing Receipt with changes noted in red.
- (X) Copy of the first page of the patent application as filed on December 23, 1999 and a copy of the returned postcard stamped by the U.S. Patent and Trademark Office.
- (X) Return prepaid postcard.



Daniel Hart  
Registration No. 40,637  
Attorney of Record

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TETRAGN.002A



PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant	:	Irena N. Merenkova	)
App. No.	:	09/471,703	)
Filed	:	December 23, 1999	)
For	:	ANALYSIS OF NUCLEOTIDE POLYMORPHISMS AT A SITE	)
Art Unit No.	:	1643	)
Examiner	:	Unknown	)

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Patents, Washington, D.C. 20231, on

May 5, 2000  
Daniel Hart  
(Date)  
Daniel Hart, Reg. No. 40,637

REQUEST FOR CORRECTION OF FILING RECEIPT

Assistant Commissioner for Patents  
Washington, D.C. 20231

Attn: Office of Initial Patent Examination's Customer Service Center

Dear Sir:

Applicant hereby respectfully requests that the Official Filing Receipt ("OFR") be corrected to show the title as ANALYSIS OF NUCLEOTIDE POLYMORPHISMS AT A SITE. A copy of the first page of the patent application as filed on December 23, 1999 and a copy of the returned postcard stamped by the U.S. Patent and Trademark Office are attached. A copy of the Official Filing Receipt is also attached with the changes noted in red.

Application No.: 09/471,703  
Filing Date: December 23, 1999

As the errors cited in the official filing receipt were incurred through the fault of the Patent Office, no fee is believed to be required. However, please charge our Deposit Account No. 11-1410 for any fees that may be incurred with this request. A duplicate of this letter is enclosed for this purpose.

Please forward the corrected Filing Receipt to the undersigned.

Respectfully submitted,

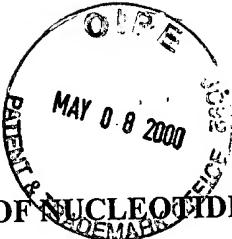
KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: May 5, 2000

By: Daniel Hart  
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TETRAGN:002A



PATENT

**ANALYSIS OF NUCLEOTIDE POLYMORPHISMS AT A SITE**

Background of the Invention

5      Field of the Invention

The present invention relates to methods of determining the identity of a polymorphic nucleotide in a target sequence having at least two variants such as a single nucleotide polymorphism, or SNP. The methods of the present invention utilize primers having sequences complementary to the region upstream of the position being analyzed.  
10 Extension of primers hybridized to target sites is carried out in the absence of a deoxyribonucleoside triphosphate (dNTP) or ribonucleoside triphosphate (rNTP) complementary to one of the polymorphic nucleotides. Differences in length between the primers and any extension products reveal the identity of the nucleotide present at the polymorphic site.

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Background of the Invention

DNA polymorphism can be due to differences in sequence or in length of a genomic region. Approximately 80% of human DNA polymorphisms are sequence polymorphisms, while only about 20% are length polymorphisms. About 90% of sequence polymorphisms are single nucleotide polymorphisms (SNPs). SNPs are genetic variations that arise from differences in the identity of a single nucleotide in a nucleic acid sequence, giving rise to two variants (sometimes called alleles) of that site. Sites having three polymorphic nucleotides have also been detected. SNPs appear to be the most widely distributed genetic markers in the human genome, occurring approximately every kilobase. Since SNPs represent the most common type of DNA sequence variation, the ability to discriminate between variants of these genetic markers is a very important tool in genetic research.  
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Many inherited diseases are the result of single point mutations at SNP sites. In some cases, the single point mutation causing nucleotide substitution in a protein-encoding gene is sufficient to actually cause the disease, as in sickle cell anemia and hemophilia. For diseases influenced by a large number of genes, including diabetes,



Date: December 23, 1999

**UTILITY/DESIGN PATENT**

(application)

Rec'd in the U.S.P.T.O. on the date stamped hereon via:

Express Mail #: EL531000980US Atty: Doh/Dop

Atty. Dkt. #: TETRAGN. 002A Applicant: I N. Merenkova.

Title: ANALYSIS OF Nucleotide Polymorphisms AT A SITE

VERIFIED BY: Asst: JEP Quality Control [Signature]

- Patent Appln. in 20 pgs. incl. Spec and 33 Claims  
 Transmittal 1 pg. 5 pgs. of Drawings  
 Preliminary Amendment in    pgs.  Power of Atty. by assignee JC688 U.S. PTO  
 Decl. and Power of Atty. copy of Assignment 09/471703  
 Decl. by Inventor(s)  Small Entity Statement(s)  
 Filed Unsigned  
 Assignment in    pgs.  
 Information Disclosure Statement; PTO-1449 w/    Ref(s)  
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12/23/99

TETRAGN.002A

PTO-103X D OH DOP  
(Rev. 6-99)

FILING RECEIPT.



UNITED STATES GOVERNMENT OF COMMERCE  
Patent and Trademark Office  
ASSISTANT SECRETARY AND COMMISSIONER  
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Washington, D.C. 20231

APPLICATION NUMBER	FILING DATE	GRP ART UNIT	FIL FEE REC'D	ATTORNEY DOCKET NO.	DRWGS	TOT CL	IND CL
09/471,703	12/23/99	1643	\$0.00	TETRAGN.002A	5	33	5

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KNOBBE MARTENS OLSON & BEAR LLP  
620 NEWPORT CENTER DRIVE  
SIXTEENTH FLOOR  
NEWPORT BEACH CA 92660



Receipt is acknowledged of this nonprovisional Patent Application. It will be considered in its order and you will be notified as to the results of the examination. Be sure to provide the U.S. APPLICATION NUMBER, FILING DATE, NAME OF APPLICANT, and TITLE OF INVENTION when inquiring about this application. Fees transmitted by check or draft are subject to collection. Please verify the accuracy of the data presented on this receipt. If an error is noted on this Filing Receipt, please write to the Office of Initial Patent Examination's Customer Service Center. Please provide a copy of this Filing Receipt with the changes noted thereon. If you received a "Notice to File Missing Parts of Application" ("Missing Parts Notice") in this application, please submit any corrections to this Filing Receipt with your reply to the "Missing Parts Notice." When the PTO processes the reply to the "Missing Parts Notice," the PTO will generate another Filing Receipt incorporating the requested corrections (if appropriate).

Applicant(s) I N MERENKOVA.

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TITLE POLYMORPHISMS  
ANALYSIS OF NUCLEOTIDE POLYMORPHISMS AT A SITE

PRELIMINARY CLASS: 435

NO DATES DOCKETED <sup>NFM</sup>  
ATTY RESPONSIBLE

DATA ENTRY BY: DIXON, DOROTHY L. TEAM: 04 DATE: 02/11/00



(See reverse for new important information)